

wherein the oligonucleotide includes at least 8 nucleotides wherein C is unmethylated and wherein X₁ and X₂ are nucleotides, wherein the cytokine is a peptide, whereby an antigen is optionally additionally administered, and wherein the antigen and the CpG oligonucleotide are not conjugated.

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22. (New) The method of claim 21, wherein the immunopotentiating cytokine is an antigen-cytokine fusion protein.

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23. (New) The method of claim 21, wherein the antigen is selected from the group consisting of a tumor antigen, a microbial antigen, and an allergen.

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24. (New) The method of claim 23, wherein the antigen is a tumor antigen.

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25. (New) The method of claim 21, wherein the antigen is administered to the subject in conjunction with the immunostimulatory CpG oligonucleotide and the immunopotentiating cytokine.

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26. (New) The method of claim 21, wherein the subject is passively exposed to the antigen.

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27. (New) The method of claim 21, wherein the subject has a neoplastic disorder.

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28. (New) The method of claim 21, wherein the subject has a viral infection.

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29. (New) The method of claim 21, wherein the subject is a non-human animal.

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30. (New) The method of claim 29, wherein the non-human animal is a vertebrate animal selected from the group consisting of a dog, a cat, a horse, a cow, a pig, a sheep, a goat, a chicken, and a primate.

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31. (New) A composition, comprising:

an effective amount for synergistically activating a dendritic cell of an immunostimulatory CpG oligonucleotide having a sequence including at least the following formula:

5' X₁CGX₂ 3'

wherein the oligonucleotide includes at least 8 nucleotides wherein C is unmethylated and wherein X₁ and X₂ are nucleotides; and a cytokine selected from the group consisting of IL-3, IL-5 and IL-12, wherein the cytokine is a peptide.

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33. (New) The composition of claim 31, wherein the cytokine is IL-3.

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34. (New) The composition of claim 31, further comprising an antigen and wherein the antigen and the CpG oligonucleotide are not conjugated.

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35. (New) The composition of claim 33, wherein the antigen is selected from the group consisting of a cancer antigen, a microbial antigen, and an allergen.

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36. (New) A method for activating a dendritic cell, comprising:

contacting a dendritic cell exposed to an antigen with an effective amount for synergistically

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activating a dendritic cell of an immunopotentiating cytokine selected from the group consisting of IL-3, IL-5 and IL-12, and an immunostimulatory CpG oligonucleotide having a sequence including at least the following formula:

5' X₁CGX₂ 3'

wherein the oligonucleotide includes at least 8 nucleotides wherein C is unmethylated and wherein X₁ and X₂ are nucleotides, wherein the cytokine is a peptide, whereby an antigen is optionally additionally administered, and wherein the antigen and the CpG oligonucleotide are not conjugated.

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37. (New) The method of claim 35, wherein the antigen is a tumor antigen.

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38. (New) A method for treating a subject having a neoplastic disorder, comprising:
administering to the tumor of a subject having a neoplastic disorder an immunopotentiating cytokine selected from the group consisting of IL-3, IL-5 and IL-12, and an immunostimulatory CpG oligonucleotide having a sequence including at least the following formula:

5' X₁CGX₂ 3'

wherein the oligonucleotide includes at least 8 nucleotides wherein C is unmethylated and wherein X₁ and X₂ are nucleotides, in an amount effective for synergistically increasing survival time of the subject with respect to a subject administered the immunostimulatory CpG oligonucleotide or the immunopotentiating cytokine alone, wherein the cytokine is a peptide.

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39. (New) The method of claim 37, wherein the tumor is selected from the group consisting of a lymphoma and a tumor of the brain, lung, ovary, breast, prostate, colon, and skin.